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Application No. 10/602,190

Amendment and Response to Office Action

*AMENDMENTS TO THE CLAIMS*

This listing of claims replaces all prior versions, and listings, of claims in the application.

1. (Currently Amended) A pharmaceutical composition, ~~comprised of~~ comprising a non-steroidal anti-inflammatory and an opiate analgesic in combination with colloidal silicate dioxide, sodium glycolate ~~glieolate~~, lactose, microcrystalline cellulose, and magnesium stearate ~~estearate and other recipients if necessary~~.
2. (Original) A pharmaceutical composition according to claim 1, wherein the non-steroidal anti-inflammatory is ketorolac tromethamine and the opiate analgesic is tramadol hydrochloride.
3. (Currently Amended) A pharmaceutical composition according to claim ~~1~~2, wherein the ketorolac tromethamine is present in the composition in a range of 0.0010 g to 0.1000 g and the tramadol hydrochloride is present in a range of 0.0010 g to 0.2000 g.
4. (Currently Amended) A pharmaceutical composition according to claim ~~1~~2, wherein the ketorolac ~~ketorolac~~ tromethamine is present in the composition in a range of 0.0010 g to 0.10000 g and the tramadol hydrochloride is present at a range of 0.0010 g to 0.2000 g, the colloidal silicate dioxide is present in a range of 0.00010g to 0.02000g, the sodium glycolate ~~glieolate~~ starch is present in a range of 0.0010g to 0.20000g, the lactose is present in the range of 0.0100g to 0.50000g, the microcrystalline cellulose is present in a range of 0.0100g to 0.50000g, the magnesium stearate is present at a proportion of 0.0001g to 0.02000g, ~~other recipients~~ and the excipient is ~~can be~~ present at a proportion of 0.0001g to 1.000g.
5. (Original) A pharmaceutical composition according to claim 1, wherein the composition it is in a capsule form.
6. (Currently Amended) A method for preparing ~~procedure for the elaboration of~~ a composition according to claim ~~1~~2, wherein it is ~~comprised of~~ comprising the following steps:

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a) ~~Mix~~ mixing the ketorolac tromethamine, the colloidal silicate dioxide, the tramadol hydrochloride, sodium glycolate ~~glycolate-starch~~, the lactose, the microcrystalline cellulose, the and magnesium stearate to produce a powdered mixture and ~~other recipients if necessary;~~

b) ~~analyze~~ analyzing the powdered ~~mix~~ mixture; and

c) ~~proceed to encapsulating and conditioning the mix~~ mixture.

7. (Currently Amended) ~~The use of A method of treating pain, the method comprising administering an effective amount of the composition according to any one of claims 1 to 6, for the pain treatment.~~